## **AMENDMENTS TO THE CLAIMS**

- 1. (Original) Process for preparing a coated metal sheet coil comprising the following steps:
  - (1) decoiling of the coiled metal sheet;
  - (2) coating the metal sheet with a curable composition comprising an (meth)acrylated oligomer which is the reaction product of a carboxyl functionalized polybutadiene comprising x equivalents of –COOH groups with (i) y equivalents of one or more (meth)acrylated monoepoxides or (ii) a mixture of z equivalents of one or more polyepoxides and at least (z-x) equivalents of an  $\alpha,\beta$ -unsaturated carboxylic acid; with z>x and  $y\ge x$ ;
  - (3) curing the composition; and
  - (4) recoiling the coated metal sheet.
- **2.** (Original) Process according to claim 1, wherein the carboxyl functionalized polybutadiene is the reaction product of a hydroxyl-terminated polybutadiene with a cyclic anhydride responding to the general formula (I):

$$\bigvee_{\mathsf{R} = \mathsf{O}}^{\mathsf{O}} \mathsf{O} \qquad \qquad \mathsf{(I)}$$

wherein R represents arylene, cycloalkylene, alkylene or alkenylene group, it being possible for R to bear alkyl, alkenyl groups, a -COOH group and/or another anhydride group.

- **3.** (Original) Process according to claim 2, wherein the anhydride is phthalic anhydride or dodecenylsuccinic anhydride.
- 4. (Currently Amended) Process according to any of claims 1 to 3 claim 1, wherein the (meth)acrylated oligomer is the reaction product of a carboxyl functionalized

polybutadiene comprising x equivalents of -COOH groups with y equivalents of one or more (meth)acrylated mono-epoxides, y being equal to x.

- **5. (Currently Amended)** Process according to any of claims 1 to 4 claim 1, wherein the (meth)acrylated mono-epoxide is chosen from glycidylacrylate and glycidylmethacrylate.
- 6. (Currently Amended) Process according to any of claims 1 to 3 claim 1, wherein the (meth)acrylated oligomer is the reaction is the reaction product of a carboxyl functionalized polybutadiene comprising x equivalents of –COOH groups with z equivalents of at least one polyepoxide and (z-x) equivalents of at least one  $\alpha,\beta$ -unsaturated carboxylic acid.
- 7. (Original) Process according to claim 6, wherein z is greater than 2x.
- 8. (Currently Amended) Process according to claim 6 or 7, wherein  $\alpha,\beta$ -unsaturated carboxylic acid is chosen from acrylic and methacrylic acid.
- 9. (Currently Amended) Process according to any of claims 1 to 3 or 6 to 8 claim 1, wherein the polyepoxide is chosen from diglycidylethers of aromatic or aliphatic diols or cycloaliphatic diepoxides.
- 10. (Original) Process according to claim 9, wherein the polyepoxide is chosen from diglycidyl ether of bisphenol-A, diglycidylether of poly(ethylene oxide-co-propylene oxide), diglycidylether of polypropylene oxide and diglycidylether of butanediol.
- 11. (Currently Amended) Process according to any of claims 1 to 3 or 6 to 10 claim 1, wherein the (meth)acrylated oligomer is prepared by adding the  $\alpha,\beta$  unsaturated carboxylic acid to the carboxyl functionalized polybutadiene before or at the latest at the same time as the polyepoxide.

- 12. (Currently Amended) Process according to any of claims 1 to 11 claim 1 wherein the (meth)acrylated oligomer is obtained by the reaction of the carboxyl functionalised polybutadiene and the mono- or polyepoxide in the presence of at least one non reactive diluent chosen from mono- or polyfunctional (meth)acrylate monomers.
- 13. (Original) Process according to claim 12, wherein the non reactive diluent is chosen from phenoxyethyl acrylate, isobornyl acrylate, n-butyl acryloyloxy ethyl carbamate and their mixtures.
- 14. (Currently Amended) Process according to any of claims 1 to 13 claim 1, wherein the curable composition comprises:
  - from 8 % to 50 % by weight of (meth)acrylated oligomer,
  - from 0 to 65 % by weight of non-reactive diluent,
  - from 0 to 60 % by weight of additional diluent chosen from copolymerizable ethylenically unsaturated monomers,
  - from 0.01 to 60 % by weight of (meth)acrylated polyepoxide,
  - from 0.01 to 5 % by weight of photoinitiator or chemical initiator, and
  - from 0 to 20 % by weight of adhesion promoter.
- 15. (Currently Amended) Process according to any of claims 1 to 14 claim 1, wherein the curing is done by electron beam or UV-radiation.